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FILE 'HOME' ENTERED AT 16:57:33 ON 03 JAN 2007

=>
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THIS COMMAND NOT AVAILABLE IN THE CURRENT FILE
Do you want to switch to the Registry File?
Choice (Y/n):
Switching to the Registry File...

Some commands only work in certain files. For example, the EXPAND command can only be used to look at the index in a file which has an index. Enter "HELP COMMANDS" at an arrow prompt (=>) for a list of commands which can be used in this file.

=> FILE REGISTRY

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 0.21 0.21

FULL ESTIMATED COST

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Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 2 JAN 2007 HIGHEST RN 916646-22-5 DICTIONARY FILE UPDATES: 2 JAN 2007 HIGHEST RN 916646-22-5

New CAS Information Use Policies, enter HELP USAGETERMS for details.

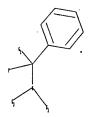
TSCA INFORMATION NOW CURRENT THROUGH June 30, 2006

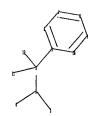
Please note that search-term pricing does apply when conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/ONLINE/UG/regprops.html

Uploading C:\Program Files\Stnexp\Queries\10537437fulorinator.str





```
chain nodes :
1 2 3 4 11 13
ring nodes :
5 6 7 8 9 10
chain bonds :
1-2 1-3 1-4 2-5 2-11 2-13
ring bonds :
5-6 5-10 6-7 7-8 8-9 9-10
exact/norm bonds :
1-2 1-3 1-4 2-13
exact bonds :
2-5 2-11
normalized bonds :
5-6 5-10 6-7 7-8 8-9 9-10
```

G1:H,C1,Br,F,I

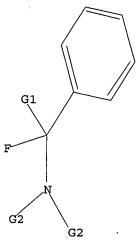
G2:CH3,CH2,n-Pr,i-Pr,n-Bu,i-Bu,s-Bu,t-Bu,Ph

Match level:
1:CLASS .2:CLASS 3:CLASS 4:CLASS 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom 11:CLASS 13:CLASS

L1 STRUCTURE UPLOADED

STR

=> d l1 L1 HAS NO ANSWERS L1



G1 H,Cl,Br,F,I G2 Me,CH2,n-Pr,i-Pr,n-Bu,i-Bu,s-Bu,t-Bu,Ph

Structure attributes must be viewed using STN Express query preparation.

=> s l1 sss full FULL SEARCH INITIATED 16:58:32 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 4035 TO ITERATE

100.0% PROCESSED 4035 ITERATIONS SEARCH TIME: 00.00.01

21 ANSWERS

L2 21 SEA SSS FUL L1

=> file caplus COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 172.10 172.31

FULL ESTIMATED COST

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of this information, without the prior written consent of CAS, is strictly prohibited. FILE COVERS 1907 - 3 Jan 2007 VOL 146 ISS 2 FILE LAST UPDATED: 2 Jan 2007 (20070102/ED) Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at: http://www.cas.org/infopolicy.html => s 12 and (?saccharide or sugar or glucose or fructose or galactose or ribose or deoxyribose or starch or chitin or cellulose) 32 L2 166701 ?SACCHARIDE 257156 SUGAR 414493 GLUCOSE 63693 FRUCTOSE 56624 GALACTOSE 27347 RIBOSE 4227 DEOXYRIBOSE 161779 STARCH 15969 CHITIN 347856 CELLULOSE L3 3 L2 AND (?SACCHARIDE OR SUGAR OR GLUCOSE OR FRUCTOSE OR GALACTOSE OR RIBOSE OR DEOXYRIBOSE OR STARCH OR CHITIN OR CELLULOSE) => d 13 1-3 ti abs bib L_3 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN De novo asymmetric syntheses of C-4-substituted sugars via an iterative . TT dihydroxylation strategy A short and highly efficient route to various C-4 substituted AB sugar lactones has been developed. The key to the overall transformation is the sequential osmium-catalyzed dihydroxylation reaction of substituted 2,4-dienoates and an allylic substitution at the C-4 position. When the Sharpless AD-mix procedure is used in a matched sense for the second dihydroxylation reaction, it results in an exceedingly diastereo- and enantioselective synthesis of several C-4-substituted sugars. AN2006:548970 CAPLUS DN 145:211276 De novo asymmetric syntheses of C-4-substituted sugars via an iterative TI dihydroxylation strategy Ahmed, Md. Moinuddin; O'Doherty, George A. ΑU CS Department of Chemistry, West Virginia University, Morgantown, WV, 26506, Carbohydrate Research (2006), 341(10), 1505-1521 SO CODEN: CRBRAT; ISSN: 0008-6215 PBElsevier B.V. DTJournal LA English os CASREACT 145:211276 RE.CNT 26 THERE ARE 26 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L3 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Deoxyfluorination of alcohols using N,N-diethyl- α , α -difluoro-(m-methylbenzyl)amine
- AB Deoxyfluorination of alcs. was carried out using N,N-diethyl- α,α -difluoro-(m-methylbenzyl)amine (DFMBA). Primary alcs. were effectively converted to fluorides under microwave irradiation or conventional heating. Deoxyfluorination of an anomeric hydroxy group in sugars by DFMBA proceeded at below room temperature and glycosyl fluorides could

be obtained in good yields. The deoxyfluorination reaction chemoselectively proceeded and various protecting groups on the sugar can survive under the reaction conditions.

AN 2004:581849 CAPLUS

DN 141:260951

TI Deoxyfluorination of alcohols using N,N-diethyl- α , α -difluoro-(m-methylbenzyl)amine

AU Kobayashi, Shingo; Yoneda, Atushi; Fukuhara, Tsuyoshi; Hara, Shoji

CS Division of Molecular Chemistry, Graduate School of Engineering, Hokkaido University, Sapporo, 060-8628, Japan

SO Tetrahedron (2004), 60(32), 6923-6930 CODEN: TETRAB; ISSN: 0040-4020

PB Elsevier Science B.V.

DT Journal

LA English

OS CASREACT 141:260951

RE.CNT 34 THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L3 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Method of fluorination using N,N-diethyl- α , α -difluorobenzylamines
- AB Disclosed is a method in which a glucide, examples of which include a monosaccharide, an oligosaccharide, a polysaccharide, a composite saccharide comprising any of these saccharides and a protein or lipid bonded thereto, a polyalc., an aldehyde, ketone, or acid of a polyalc., a derivative or condensate of any of these, is reacted with a fluorinating agent represented by the general formula of RCF2-Y(R1)R2 [y = N, P; R-R2 are same or different group selected from H and each (un) substituted alkyl and aryl; or ≥2 of R-R2 groups are bonded to each other to form a ring] either thermally or by irradiation with microwave or an electromagnetic wave with a wavelength around the microwave region. By the method, fluorination reaction can be safely conducted position-selectively even in a temperature range of 150 to 200°, in which fluorination has conventionally been difficult. method in which the reactants are irradiated with microwave or an electromagnetic wave with a wavelength around the microwave region is applicable to substrates other than glucides. When a complex compound comprising HF and a base, for example, is reacted with a substrate by irradiation with microwave, fluorination in a specific position which has been difficult in conventional techniques proceeds highly selectively in a short time efficiently and safely. Thus, 10 mmol Me 2,3-O-isopropylidene- β -D-ribofuranoside, 12 mmol N, N-diethyl- α , α -difluoro-3methylbenzylamine, and 20 mL heptane were added to a glass vessel reaction vessel coated with fluorinated resin, heated with 100° with stirring, and allowed to react for 50 min to give 55% Me 2,3-0-isopropylidene-5-deoxy-5-fluoro-β-D-ribofuranoside.

AN 2004:493719 CAPLUS

DN 141:38808

TI Method of fluorination using N,N-diethyl- α , α -difluorobenzylamines

IN Hara, Shoji; Fukuhara, Tsuyoshi

PA Mitsubishi Gas Chemical Company, Inc., Japan

SO PCT Int. Appl., 50 pp.

CODEN: PIXXD2

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004050676	A1	20040617	WO 2003-JP15336	20031201
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	JP 2004182664		, RO, SE, S 20040702	I, SK, TR JP 2002-352968	20021204

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A 20040708 JP 2002-358249
A1 20050831 EP 2003-775984
      JP 2004189655
      EP 1568703
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 PRAI JP 2002-352968
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      CASREACT 141:38808; MARPAT 141:38808
 RE.CNT 19
            THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD
               ALL CITATIONS AVAILABLE IN THE RE FORMAT
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         166709 ?ACCHARIDE
          49821 FLUORINAT?
 L4
            483 (SUGAR OR ?ACCHARIDE) AND (FLUORINAT?)
 => d l4 1-5 ti
      ANSWER 1 OF 483 CAPLUS COPYRIGHT 2007 ACS on STN
 TI
      Design, synthesis, and biological evaluation of novel iso-D-2',3'-dideoxy-
      3'-fluorothianucleoside derivatives
      ANSWER 2 OF 483 CAPLUS COPYRIGHT 2007 ACS on STN
 L4
 ТT
      Preparation of highly fluorinated carboxylic acids and their
      application as protective groups in fluorous synthesis
 L4
      ANSWER 3 OF 483 CAPLUS COPYRIGHT 2007 ACS on STN
      Fluorinated analogues of biological molecules: accessing new
 TI
      chemical, physical and biological properties
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     ANSWER 4 OF 483 CAPLUS COPYRIGHT 2007 ACS on STN
 TI
     A synthesis of 2-fluoroglucal derivatives
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      ANSWER 5 OF 483 CAPLUS COPYRIGHT 2007 ACS on STN
 TI
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NEWS
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                 LOGOFF HOLD duration extended to 120 minutes
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                 E-mail format enhanced
NEWS 13
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NEWS 14
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NEWS 15
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NEWS 16
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NEWS 17
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NEWS 27
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NEWS EXPRESS
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              AND CURRENT DISCOVER FILE IS DATED 25 SEPTEMBER 2006.
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              Welcome Banner and News Items
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